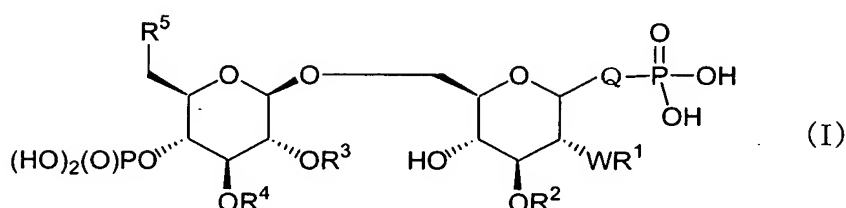


## CLAIMS

1. A compound represented by the following general formula:



wherein Q represents an oxygen atom, a C<sub>1</sub>-C<sub>3</sub> alkylene group, a -O-Alk- group or a -O-Alk-O- group (in which Alk represents a C<sub>1</sub>-C<sub>3</sub> alkylene group),

W represents an oxygen atom or a -NH- group,

R<sup>1</sup> (when W is a -NH- group) represents a C<sub>1</sub>-C<sub>20</sub> alkanoyl group which may be substituted by at least one group selected from the following Substituent group A, a C<sub>3</sub>-C<sub>20</sub> alkenoyl group which may be substituted by at least one group selected from the following Substituent group A or a C<sub>3</sub>-C<sub>20</sub> alkynoyl group which may be substituted by at least one group selected from the following Substituent group A,

R<sup>1</sup> (when W is an oxygen atom), R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup>, which may be the same or different, represent a hydrogen atom, a C<sub>1</sub>-C<sub>20</sub> alkyl group which may be substituted by at least one group selected from the following Substituent group A, a C<sub>2</sub>-C<sub>20</sub> alkenyl group which may be substituted by at least one group selected from the following Substituent group A, a C<sub>2</sub>-C<sub>20</sub> alkynyl group which may be substituted by at least one group selected from the following Substituent group A, a C<sub>1</sub>-C<sub>20</sub> alkanoyl group which may be substituted by at least one group selected from the following Substituent group A, a C<sub>3</sub>-C<sub>20</sub> alkenoyl group which may be substituted by at least one group selected from the following Substituent group A or a C<sub>3</sub>-C<sub>20</sub> alkynoyl

group which may be substituted by at least one group selected from the following Substituent group A,

$R^5$  represents a hydrogen atom, a halogen atom, a hydroxyl group, a  $C_1$ - $C_6$  alkoxy group which may have an oxo group, a  $C_2$ - $C_6$  alkenyloxy group which may have an oxo group or a  $C_2$ - $C_6$  alkynyloxy group which may have an oxo group,

the Substituent group A consisting of a halogen atom, a hydroxyl group, an oxo group, a  $C_1$ - $C_{20}$  alkoxy group which may have an oxo group, a ( $C_1$ - $C_{20}$  alkoxy)  $C_1$ - $C_{20}$  alkoxy group, a {( $C_1$ - $C_{20}$  alkoxy)  $C_1$ - $C_{20}$  alkoxy}  $C_1$ - $C_{20}$  alkoxy group, a  $C_2$ - $C_{20}$  alkenyloxy group which may have an oxo group, a  $C_2$ - $C_{20}$  alkynyloxy group which may have an oxo group, a  $C_1$ - $C_{20}$  alkanoyloxy group which may have an oxo group, a  $C_3$ - $C_{20}$  alkenoyloxy group which may have an oxo group and a  $C_3$ - $C_{20}$  alkynoyloxy group which may have an oxo group, or a pharmacologically acceptable salt thereof.

2. The compound according to claim 1, wherein W is a -NH- group and  $R^1$  is a  $C_8$ - $C_{18}$  alkanoyl or  $C_8$ - $C_{18}$  alkenoyl group, which may have a substituent selected from the Substituent group A, or a pharmacologically acceptable salt thereof.

3. The compound according to claim 1, wherein W is a -NH- group and  $R^1$  is a  $C_{10}$ - $C_{18}$  alkanoyl or  $C_{10}$ - $C_{18}$  alkenoyl group, which may have a substituent selected from the Substituent group A, or a pharmacologically acceptable salt thereof.

4. The compound according to claim 1, wherein W is a -NH- group and  $R^1$  is a  $C_{12}$ - $C_{16}$  alkanoyl or  $C_{12}$ - $C_{16}$  alkenoyl group, which may have a substituent selected from the Substituent group A, or a pharmacologically acceptable salt thereof.

5. The compound according to claim 1, wherein W is an oxygen atom and  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$ , which may be the same or different, are a  $C_4$ - $C_{18}$  alkyl,  $C_4$ - $C_{18}$  alkenyl,  $C_4$ - $C_{18}$  alkanoyl or  $C_4$ - $C_{18}$  alkenoyl group, which may have a substituent selected from the Substituent group A, or a pharmacologically acceptable salt thereof.
6. The compound according to claim 1, wherein W is an oxygen atom and  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$ , which may be the same or different, are a  $C_8$ - $C_{18}$  alkyl,  $C_8$ - $C_{18}$  alkenyl,  $C_8$ - $C_{18}$  alkanoyl or  $C_8$ - $C_{18}$  alkenoyl group, which may have a substituent selected from the Substituent group A, or a pharmacologically acceptable salt thereof.
7. The compound according to claim 1, wherein W is an oxygen atom and  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$ , which may be the same or different, are a  $C_{10}$ - $C_{18}$  alkyl,  $C_{10}$ - $C_{18}$  alkenyl,  $C_{10}$ - $C_{18}$  alkanoyl or  $C_{10}$ - $C_{18}$  alkenoyl group, which may have a substituent selected from the Substituent group A, or a pharmacologically acceptable salt thereof.
8. The compound according to claim 1, wherein W is an oxygen atom and  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$ , which may be the same or different, are a  $C_{12}$ - $C_{16}$  alkyl,  $C_{12}$ - $C_{16}$  alkenyl,  $C_{12}$ - $C_{16}$  alkanoyl or  $C_{12}$ - $C_{16}$  alkenoyl group, which may have a substituent selected from the Substituent group A, or a pharmacologically acceptable salt thereof.
9. The compound according to claim 1, wherein W is an oxygen atom,  $R^1$  and  $R^3$ , which may be the same or different, are a  $C_{12}$ - $C_{16}$  alkanoyl or  $C_{12}$ - $C_{16}$  alkenoyl group, which may have a substituent selected from the Substituent group A, and  $R^2$  and  $R^4$ , which may be the same or different, are a  $C_{12}$ - $C_{16}$  alkyl or a  $C_{12}$ - $C_{16}$  alkenyl group, which may have a substituent selected from the Substituent group A, or a pharmacologically acceptable salt thereof.

10. The compound according to claim 1, wherein W is an oxygen atom,  $R^1$  and  $R^3$ , which may be the same or different, are a decanoyl, dodecanoyl, tetradecanoyl, dodecenoyl, tetradecenoyl or octadecenoyl group, which may have a substituent selected from the Substituent group A, and  $R^2$  and  $R^4$ , which may be the same or different, are decyl, dodecyl, tetradecyl, dodecenyl, tetradecenyl or octadecenyl group, which may have a substituent selected from the Substituent group A, or a pharmacologically acceptable salt thereof.

11. The compound according to any one of claims 1 to 10, wherein the substituent selected from the Substituent group A is a fluorine atom, a hydroxyl group, a  $C_1$ - $C_{20}$  alkoxy group, a  $C_{12}$ - $C_{14}$  alkenyloxy group, a  $C_{12}$ - $C_{14}$  alkanoyloxy group or a  $C_{12}$ - $C_{14}$  alkenoyloxy group, or a pharmacologically acceptable salt thereof.

12. The compound according to any one of claims 1 to 10, wherein the substituent selected from the Substituent group A is a dodecyloxy group, a tetradecyloxy group, a dodecenyloxy group, a tetradecenyloxy group, a dodecanoyloxy group, a tetradecanoyloxy group, a dodecenoyloxy group, a tetradecenoyloxy group or an octadecenoyl group, or a pharmacologically acceptable salt thereof.

13. The compound according to any one of claims 1 to 12, wherein  $R^5$  is a halogen atom, a hydroxyl group or an unsubstituted  $C_1$ - $C_6$  alkoxy group, or a pharmacologically acceptable salt thereof.

14. The compound according to any one of claims 1 to 12, wherein  $R^5$  is a fluorine atom, a hydroxyl group or a methoxy group, or a pharmacologically acceptable salt thereof.

15. The compound according to any one of claims 1 to 14, wherein Q is an oxygen atom, or a pharmacologically acceptable salt thereof.
16. The compound according to any one of claims 1 to 14, wherein Q is a phosphonoethyl group, or a pharmacologically acceptable salt thereof.
17. The compound according to any one of claims 1 to 16, wherein position 1 of the right-side glucose or glucosamine takes the  $\alpha$  configuration, or a pharmacologically acceptable salt thereof.
18. Phosphono 3-O-decyl-2-deoxy-6-O-{3-O-[(R)-3-methoxydecyl]-6-O-methyl-2-O-[(Z)-11-octadecenoyl]-4-O-phosphono- $\beta$ -D-glucopyranosyl}-2-(3-oxotetradecanoylamino)- $\alpha$ -D-glucopyranoside,  
 phosphono 3-O-decyl-2-deoxy-6-O-{3-O-[(R)-3-methoxydecyl]-6-O-methyl-2-O-[(Z)-11-octadecenyl]-4-O-phosphono- $\beta$ -D-glucopyranosyl}-2-(3-oxotetradecanoylamino)- $\alpha$ -D-glucopyranoside,  
 phosphono 3-O-decyl-2-deoxy-6-O-{3-O-[(R)-3-methoxydecyl]-2-O-[(Z)-11-octadecenoyl]-4-O-phosphono- $\beta$ -D-glucopyranosyl}-2-(3-oxotetradecanoylamino)- $\alpha$ -D-glucopyranoside,  
 phosphono 3-O-decyl-2-deoxy-6-O-{3-O-[(R)-3-methoxydecyl]-2-O-[(Z)-11-octadecenyl]-4-O-phosphono- $\beta$ -D-glucopyranosyl}-2-(3-oxotetradecanoylamino)- $\alpha$ -D-glucopyranoside,  
 2-(phosphonooxy)ethyl 3-O-decyl-2-deoxy-6-O-{3-O-[(R)-3-methoxydecyl]-6-O-methyl-2-O-[(Z)-11-octadecenoyl]-4-O-phosphono- $\beta$ -D-glucopyranosyl}-2-(3-oxotetradecanoylamino)- $\alpha$ -D-glucopyranoside,

2-(phosphonooxy)ethyl 3-O-decyl-2-deoxy-6-O-{3-O-[(R)-3-methoxydecyl]-2-O-[(Z)-11-octadecenoyl]-4-O-phosphono- $\beta$ -D-glucopyranosyl}-2-(3-oxotetradecanoylamino)- $\alpha$ -D-glucopyranoside,

2-(phosphonooxy)ethyl 6-O-{3-O-[(R)-3-methoxydecyl]-6-O-methyl-2-O-[(Z)-11-octadecenoyl]-4-O-phosphono- $\beta$ -D-glucopyranosyl}-2,3-di-O-dodecyl- $\alpha$ -D-glucopyranoside,

phosphono 3-O-decyl-6-O-{3-O-[(R)-3-methoxydecyl]-6-O-methyl-2-O-[(Z)-11-octadecenoyl]-4-O-phosphono- $\beta$ -D-glucopyranosyl}-2-O-(3-oxotetradecanoyl)- $\alpha$ -D-glucopyranoside,

phosphono 3-O-decyl-6-O-{3-O-[(R)-3-methoxydecyl]-2-O-[(Z)-11-octadecenoyl]-4-O-phosphono- $\beta$ -D-glucopyranosyl}-2-O-(3-oxotetradecanoyl)- $\alpha$ -D-glucopyranoside,

2-(phosphonooxy)ethyl 2,3-di-O-dodecyl-6-O-{6-O-methyl-3-O-[(R)-3-methoxydecyl]-2-O-[(Z)-11-octadecenoyl]-4-O-phosphono- $\beta$ -D-glucopyranosyl}- $\alpha$ -D-glucopyranoside or

phosphono 6-O-{4-O-phosphono-3-O-[(R)-3-methoxydecyl]-6-O-methyl-2-O-[(Z)-11-octadecenoyl]- $\beta$ -D-glucopyranosyl}-3-O-dodecyl-2-O-[(R)-3-hydroxytetradecyl]- $\alpha,\beta$ -D-glucopyranoside according to claim 1, or a pharmacologically acceptable salt thereof.

19. Phosphono 3-O-decyl-6-O-{3-O-[(R)-3-methoxydecyl]-6-O-methyl-2-O-[(Z)-11-octadecenoyl]-4-O-phosphono- $\beta$ -D-glucopyranosyl}-2-O-(3-oxotetradecanoyl)- $\alpha$ -D-glucopyranoside or

phosphono 3-O-decyl-6-O-{3-O-[(R)-3-methoxydecyl]-2-O-[(Z)-11-octadecenoyl]-4-O-phosphono- $\beta$ -D-glucopyranosyl}-2-O-(3-oxotetradecanoyl)- $\alpha$ -D-glucopyranoside according to claim 1, or a pharmacologically acceptable salt thereof.

20. A medicament comprising the compound according to any one of claims 1 to 19 as an active ingredient.
21. An agent for prophylaxis or treatment of inflammation, comprising the compound according to any one of claims 1 to 19 as an active ingredient.
22. An agent for prophylaxis or treatment of an autoimmune disease, comprising the compound according to any one of claims 1 to 19 as an active ingredient.
23. An agent for prophylaxis or treatment of sepsis, comprising the compound according to any one of claims 1 to 19 as an active ingredient.
24. An immunosuppressive agent comprising the compound according to any one of claims 1 to 19 as an active ingredient.
25. A prognosis-improving agent after coronary artery bypass surgery, comprising the compound according to any one of claims 1 to 19 as an active ingredient.
26. Use of the compound according to any one of claims 1 to 19 or a pharmacologically acceptable salt thereof for producing a pharmaceutical composition.
27. A method for prophylaxis or treatment of inflammation, which comprises administering a pharmacologically effective dose of the compound according to any one of claims 1 to 19 or a pharmacologically acceptable salt thereof to a warm-blooded animal.

28. A method for prophylaxis or treatment of an autoimmune disease, which comprises administering a pharmacologically effective dose of the compound according to any one of claims 1 to 19 or a pharmacologically acceptable salt thereof to a warm-blooded animal.

29. A method for prophylaxis or treatment of sepsis, which comprises administering a pharmacologically effective dose of the compound according to any one of claims 1 to 19 or a pharmacologically acceptable salt thereof to a warm-blooded animal.

30. A method for immunosuppression, which comprises administering a pharmacologically effective dose of the compound according to any one of claims 1 to 19 or a pharmacologically acceptable salt thereof to a warm-blooded animal.

31. A method for improving prognosis after coronary artery bypass surgery, which comprises administering a pharmacologically effective dose of the compound according to any one of claims 1 to 19 or a pharmacologically acceptable salt thereof to a warm-blooded animal.